

HIV

Unilateral syphilitic peri optic neuritis in a patient coinfectd with human immunodeficiency virus type 1

Medhat S T Basta, K Nathan Sankar, Margaret Dayan

Sex Transm Infect 2007;**83**:183–184. doi: 10.1136/sti.2006.024067

Peri optic neuritis caused by secondary syphilis is a rare ophthalmic manifestation in the HIV-infected host. Early diagnosis and treatment of this condition is required to prevent further visual damage. We report a case of unilateral syphilitic peri optic neuritis in a patient coinfectd with HIV-1.

Peri optic neuritis usually occurs as a bilateral condition in secondary syphilis. To our knowledge, this report is the first that describes the rare occurrence of unilateral syphilitic peri optic neuritis in a patient who is also infected with HIV-1. The diagnosis of this condition in this patient influenced the management of secondary syphilis and prevented progression of inflammation to the optic nerve substance and permanent visual loss.

CASE REPORT

A 42-year-old homosexual man presented to the genitourinary medicine department with a 1-week history of orogenital ulcers, associated with a rash on the palms, soles and trunk. Although he was seeing cobweb-like images in his right eye for the same duration, he reported this 2 days later to the ophthalmologists at the eye casualty. His medical history was unremarkable with no previous ocular symptoms. On examination, the rash was suggestive of secondary syphilis. He had bilateral non-tender inguinal lymphadenopathy. Dark ground microscopy of the genital ulcer exudate was negative for spirochaetes. The right optic disc was swollen (fig 1) with a few cells in the overlying vitreous and the left optic disc was unaffected. The visual acuities were 6/5 right and 6/18 left (an amblyopic eye). There was no relative afferent pupil defect or evidence of retinopathy in either eye. Visual fields and colour vision were normal.

Syphilis serology showed reactive enzyme immunoassay by both Abbott Murex ICE Syphilis (Abbott Murex, Dartford, UK) and Dade Behring Enzygnost (Dade Behring Enzygnost,

Marburg, Germany), IgM enzyme-linked immunosorbent assay, treponema pallidum serodia particle agglutination test at $\geq 1:1280$ and venereal diseases reference laboratory (VDRL) test at 1:256. HIV-1 antibody test was positive with HIV-1 RNA polymerase chain reaction of 25 500 copies/ml, and a CD4 count of 360 cells/ μ l. CT scan of the head was normal. Cerebrospinal fluid examination findings were: normal opening pressure; white cell count 55 cells/ml (82% lymphocytes); protein 1.2 g/l; glucose 2.9 mmol/l (plasma glucose 5.4 mmol/l); and VDRL negative and TPPA reactive at 1:160. Cerebrospinal fluid results were thus inconclusive of neurosyphilis. A diagnosis of secondary syphilis with associated peri optic neuritis in the right eye was made and the patient was treated with procaine penicillin 1.8 G intramuscularly and oral probenidol 500 mg four times a day for 17 days. At 2 months after treatment, there was complete resolution of the right optic disc swelling and a 32-fold decline in VDRL titre to 1:8. The patient has remained asymptomatic, and at the last follow-up visit at 15 months the VDRL was negative.

DISCUSSION

This is a case of unilateral optic disc swelling diagnosed in a patient with secondary syphilis who was also infected with HIV-1. The visual acuity in the affected eye remained unimpaired which, in the presence of normal intracranial pressure and inflammatory cells in the vitreous, is consistent with peri optic neuritis—that is, inflammation of the optic nerve meninges but sparing the optic nerve fibres themselves. Peri optic neuritis therefore produces optic disc swelling without affecting visual function. In addition to secondary syphilis, this condition has been described in a variety of inflammatory and infectious eye diseases including meningitis and sarcoidosis.¹

Peri optic neuritis in secondary syphilis usually occurs as a bilateral condition,^{2,3} whereas syphilitic optic neuropathy is more often unilateral. Unilateral syphilitic peri optic neuritis

Abbreviation: VDRL, venereal diseases reference laboratory

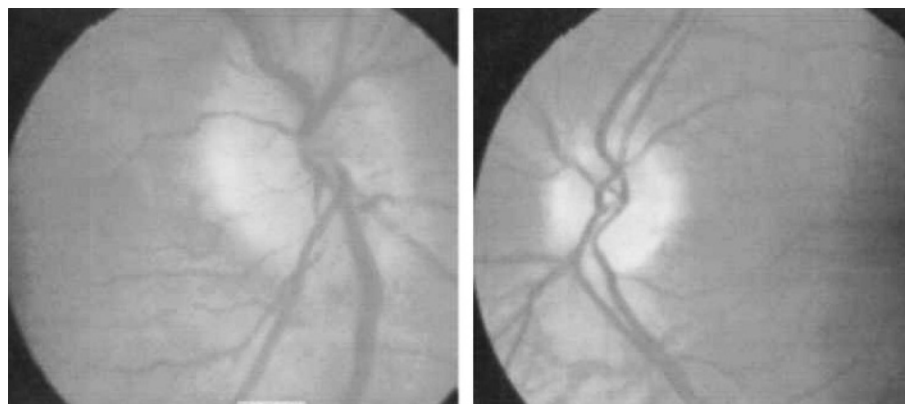


Figure 1 Right optic disc swelling in syphilitic peri optic neuritis, left optic disc is normal.

Learning points/key messages from the case report

- Syphilis should always be included in the differential diagnosis of patients presenting with optic disc swelling, especially in those at risk of sexually transmitted infections
- Patients with secondary syphilis, especially those coinfectd with HIV-1, should be questioned about visual symptoms. If they have even mild visual problems, they should have an ophthalmological review
- The diagnosis of syphilitic ocular disease, as in this case, affected the way secondary syphilis was managed.
- Syphilitic eye disease can progress more rapidly in HIV-infected patients, hence they need early diagnosis and prompt treatment.

presenting as the big blind spot syndrome has been reported once previously.⁴ Ocular syphilis is often more severe in HIV-infected patients⁵ and its occurrence does affect the way syphilis is treated.³ Early recognition and appropriate management of these cases are therefore vital to prevent progression. The ineffectiveness of conventional benzathine penicillin in the treatment of ophthalmic manifestations of secondary syphilis in the HIV-infected host has been established although a procaine penicillin-containing regimen for the treatment of neurosyphilis is adequate.³

We advocate that patients with secondary syphilis, especially those coinfectd with HIV-1, should be questioned about visual symptoms, and those who have even mild visual problems should have an ophthalmological review. Conversely, syphilis should always be included in the differential of patients

presenting with optic disc swelling, especially in those patients at risk of sexually transmitted infections. Adequate management of this case resolved the syphilitic perioptic neuritis before there was progression of inflammation to the optic nerve substance and permanent visual loss.

AUTHORS CONTRIBUTIONS

KNS and MD were involved in the initial presentation and care of the patient. MSTB and KNS continue to follow-up the patient for his HIV care. MSTB and MD carried out the literature review. MSTB wrote the initial manuscript, which KNS and MD reviewed and edited. MSTB rewrote the final manuscript for submission.

Authors' affiliations

Medhat S T Basta, K Nathan Sankar, Department of Genitourinary Medicine, Newcastle General Hospital, Newcastle upon Tyne, UK

Margaret Dayan, Department of Ophthalmology, Royal Victoria Infirmary, Newcastle upon Tyne, UK

Competing interests: None declared.

Correspondence to: Dr M S T Basta, Genitourinary medicine department, Newcastle General Hospital, Westgate Road, Newcastle upon Tyne NE4 6BE, UK; medhat.basta@newcastle-pct.nhs.uk

Accepted 11 January 2007

REFERENCES

- 1 **Walsh FB**, Hoyt WF. In: *Clinical neuro-ophthalmology*, 3rd edn, Vol.2. Baltimore: Williams & Wilkins, 1969:1582.
- 2 **Rush JA**, Ryan EJ. Syphilitic optic perineuritis. *Am J Ophthalmol* 1981;**91**:404–6.
- 3 **McLeish WM**, Pulido JS, Holland S, *et al*. The ocular manifestations of syphilis in the human immunodeficiency virus type 1-infected host. *Ophthalmology* 1990;**97**:196–203.
- 4 **McBurney J**, Rosenberg ML. Unilateral syphilitic optic perineuritis presenting as the big blind spot syndrome. *J Clin Neuroophthalmol* 1987;**7**:167–9.
- 5 **Ahmed I**, Ai E, Chang E, *et al*. *Ophthalmic manifestations of HIV. HIV Insite Knowledge Base Chapter*, 2005.